

PATENT COOPERATION TREATY
PCT
INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY
(Chapter II of the Patent Cooperation Treaty)
(PCT Article 36 and Rule 70)

REC'D 09 AUG 2005

WIPO

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Applicant's or agent's file reference 42968PCX329/29 KM	FOR FURTHER ACTION		See Form PCT/IPEA/416
International application No. PCT/NZ2004/000184	International filing date (<i>day/month/year</i>) 13 August 2004	Priority date (<i>day/month/year</i>) 15 August 2003	
International Patent Classification (IPC) or national classification and IPC Int. Cl. 7 A61K 31/407; A61P 9/10, 9/12			
Applicant AGRESEARCH LIMITED et al			

<p>1. This report is the international preliminary examination report, established by this International Preliminary Examining Authority under Article 35 and transmitted to the applicant according to Article 36.</p> <p>2. This REPORT consists of a total of 4 sheets, including this cover sheet.</p> <p>3. This report is also accompanied by ANNEXES, comprising:</p> <p>a. <input checked="" type="checkbox"/> (<i>sent to the applicant and to the International Bureau</i>) a total of 17 sheets, as follows:</p> <p><input checked="" type="checkbox"/> sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions).</p> <p><input type="checkbox"/> sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box.</p> <p>b. <input type="checkbox"/> (<i>sent to the International Bureau only</i>) a total of (indicate type and number of electronic carrier(s)) , containing a sequence listing and/or table related thereto, in computer readable form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the Administrative Instructions).</p> <p>4. This report contains indications relating to the following items:</p> <table> <tbody> <tr> <td><input checked="" type="checkbox"/></td> <td>Box No. I</td> <td>Basis of the report</td> </tr> <tr> <td><input type="checkbox"/></td> <td>Box No. II</td> <td>Priority</td> </tr> <tr> <td><input type="checkbox"/></td> <td>Box No. III</td> <td>Non-establishment of opinion with regard to novelty, inventive step and industrial applicability</td> </tr> <tr> <td><input type="checkbox"/></td> <td>Box No. IV</td> <td>Lack of unity of invention</td> </tr> <tr> <td><input checked="" type="checkbox"/></td> <td>Box No. V</td> <td>Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement</td> </tr> <tr> <td><input type="checkbox"/></td> <td>Box No. VI</td> <td>Certain documents cited</td> </tr> <tr> <td><input type="checkbox"/></td> <td>Box No. VII</td> <td>Certain defects in the international application</td> </tr> <tr> <td><input type="checkbox"/></td> <td>Box No. VIII</td> <td>Certain observations on the international application</td> </tr> </tbody> </table>	<input checked="" type="checkbox"/>	Box No. I	Basis of the report	<input type="checkbox"/>	Box No. II	Priority	<input type="checkbox"/>	Box No. III	Non-establishment of opinion with regard to novelty, inventive step and industrial applicability	<input type="checkbox"/>	Box No. IV	Lack of unity of invention	<input checked="" type="checkbox"/>	Box No. V	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement	<input type="checkbox"/>	Box No. VI	Certain documents cited	<input type="checkbox"/>	Box No. VII	Certain defects in the international application	<input type="checkbox"/>	Box No. VIII	Certain observations on the international application
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<input type="checkbox"/>	Box No. VIII	Certain observations on the international application																						

Date of submission of the demand 1 February 2005	Date of completion of the report 26 July 2005
Name and mailing address of the IPEA/AU AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRALIA E-mail address: pct@ipaaustralia.gov.au Facsimile No. (02) 6285 3929	Authorized Officer S. Chew Telephone No. (02) 6283 2248

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/NZ2004/000184

Box No. I Basis of the report

1. With regard to the language, this report is based on the international application in the language in which it was filed, unless otherwise indicated under this item.
 - This report is based on translations from the original language into the following language which is the language of a translation furnished for the purposes of:
 - international search (under Rules 12.3 and 23.1 (b))
 - publication of the international application (under Rule 12.4)
 - international preliminary examination (under Rules 55.2 and/or 55.3)
2. With regard to the elements of the international application, this report is based on (*replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report*):
 - the international application as originally filed/furnished
 - the description:
 - pages 1-33 as originally filed/furnished
 - pages* received by this Authority on with the letter of
 - pages* received by this Authority on with the letter of
 - the claims:
 - pages as originally filed/furnished
 - pages* as amended (together with any statement) under Article 19
 - pages*34-50 received by this Authority on 1 February 2005 with the letter of 1 February 2005
 - pages* received by this Authority on with the letter of
 - the drawings:
 - pages 1/5 – 5/5 as originally filed/furnished
 - pages* received by this Authority on with the letter of
 - pages* received by this Authority on with the letter of
 - a sequence listing and/or any related table(s) - see Supplemental Box Relating to Sequence Listing.
3. The amendments have resulted in the cancellation of:
 - the description, pages
 - the claims, Nos.
 - the drawings, sheets/figs
 - the sequence listing (*specify*):
 - any table(s) related to the sequence listing (*specify*):
4. This report has been established as if (some of) the amendments annexed to this report and listed below had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).
 - the description, pages
 - the claims, Nos.
 - the drawings, sheets/figs
 - the sequence listing (*specify*):
 - any table(s) related to the sequence listing (*specify*):

* If item 4 applies, some or all of those sheets may be marked "superseded."

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/NZ2004/000184

Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Claims 1-49	YES
	Claims	NO
Inventive step (IS)	Claims 1-49	YES
	Claims	NO
Industrial applicability (IA)	Claims 1-49	YES
	Claims	NO

2. Citations and explanations (Rule 70.7)

This report has considered the following documents cited in the International Search Report:

- D1 WO 2003/105868
- D2 Miles C. et al.
- D3 Munday-Finch S. et al. J. Agric. Food Chem. 1995
- D4 Munday-Finch S. et al. J. Agric. Food Chem. 1998
- D5 Munday-Finch S. et al. J. Agric. Food Chem. 1997
- D6 Derwent Abstract Accession No. 92-308267/38
- D7 Munday-Finch S. et al. J. Agric. Food Chem. 1996

NOVELTY (N), INVENTIVE STEP (IS) : Claims 1-49

D1 discloses lolitrem A, B, C, E, F, H, N, lolitrem N-31-epimer, lolitriol, lolilline, lolitriol, lolicines A and B and their use as potassium channel blockers for the treatment of ocular hypertension or glaucoma (see pages 5, 7, 13 and claim 1).

D2 discloses the isolation and structures of lolitrem B and E including their biosynthetic route from lolitriol (see abstract and figure 1).

D3 discloses the isolation of lolitrem A, its structure and structures of lolitrem B, C and E (see abstract and figure 1).

D4 discloses the isolation of lolicines A and B, lolitriol and lolitrem N and has provided evidence for 31-epitolitrem N and 31-epitolitrem F (see abstract and figure 1).

D5 discloses lolilline, lolitrem A, B, E and lolitriol (see figures 1 and 3).

D6 discloses some lolitrem derivatives used for the preparation of haptens for the production of antibodies.

D7 discloses lolitrem F, lolitrem B, 31-epitolitrem B, 31-epitolitrem F and lolitriol (see abstract, figures 1 and 4).

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

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Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

Continuation of Box No. V:

None of D1-D7 disclose or fairly suggest alone or in combination, a method of preventing repolarisation or hyperpolarisation of a cell wherein the cell contains a BK channel, comprising the administration to the cell of a composition containing a BK channel antagonist as defined in the claims, or a composition comprising a BK channel antagonist compound containing the moiety shown in structures (VII), (IX), (XII) and (XIII).

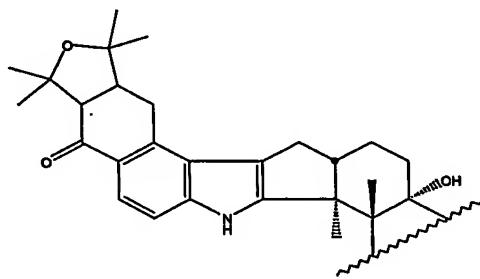
Therefore claims 1-49 are novel and have an inventive step.

INDUSTRIAL APPLICABILITY (IA): Claims 1-49

Claims 1-49 have industrial applicability.

WHAT WE CLAIM IS:

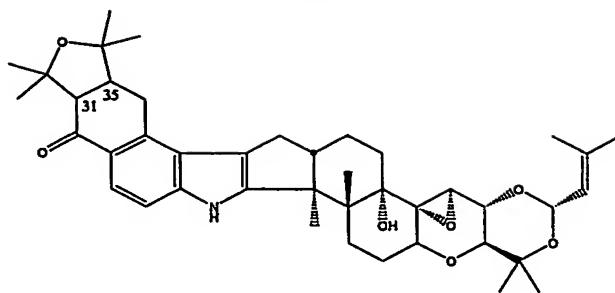
1. A method of preventing repolarisation or hyperpolarisation of a cell, wherein the cell contains a BK channel, including the administration to the cell of at least one pharmacologically effective amount of composition containing a BK channel
- 5 antagonist containing the moiety shown in structure (I):



STRUCTURE (I)

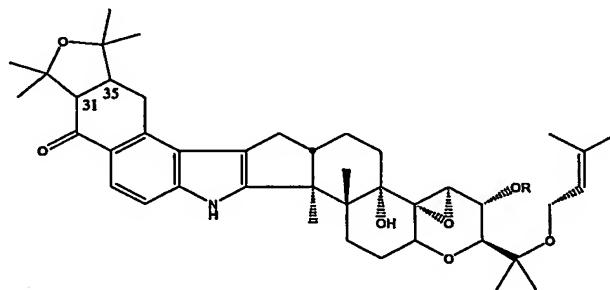
or derivatives thereof.

2. The method as claimed in claim 1 wherein the derivatives of structure (I) are selected from the group consisting of: salts, analogues, isomers, and combinations thereof.
3. The method as claimed in claim 1 or claim 2 wherein the antagonist compound is selected from the group consisting of: lolitrem B, lolitrem A, lolitrem F, 31-*epi*lolitrem F, 31-*epi*lolitrem B, lolitrem E, lolitrem E acetate, lolitrem L, lolitrem G, lolitrem C, lolitrem M, lolitriol, lolitriol acetate, lolitrem N, lolitrem J, lolitrem H, lolitrem K, lolicine A and B, 30-desoxy lolitrem B-30 α -ol, 30-desoxy-31-*epi*lolitrem B-30 α -ol, 30-desoxylolitrem B-30-ene lolilline and combinations thereof.
4. The method as claimed in claim 1 or claim 2 wherein the antagonist compound is selected from the group consisting of:



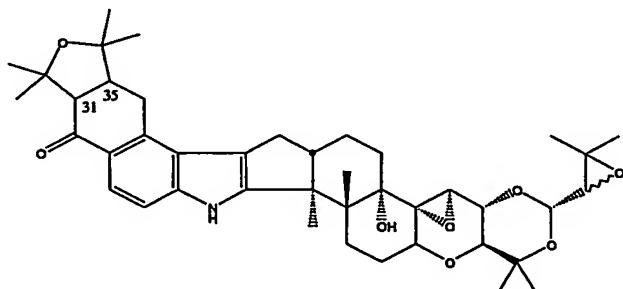
STRUCTURE (II)

- which includes compounds selected from the group consisting of: lolitrem B
= 31 α , 35 β stereochemistry; 31-epilolitrem B = 31 β , 35 β stereochemistry; lolitrem F
5 = 31 α , 35 α ; 31-epilolitrem F = 31 β , 35 α ;



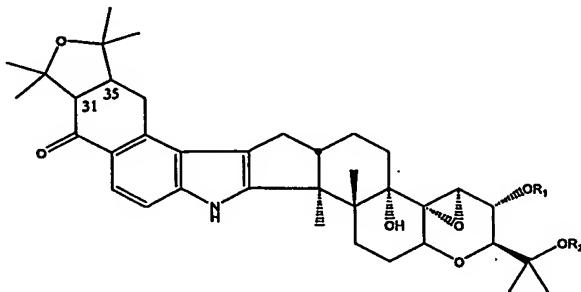
STRUCTURE (III)

- which includes compounds selected from the group consisting of: lolitrem E
= 31 α , 35 β stereochemistry where R = H or acetate; lolitrem L = 31 α , 35 α
10 stereochemistry where R = H or acetate;



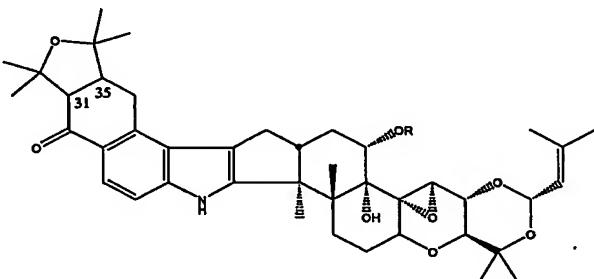
STRUCTURE (IV)

which includes compounds selected from the group consisting of: lolitrem A = $31\alpha, 35\beta$ stereochemistry; lolitrem G = $31\alpha, 35\alpha$ stereochemistry;



STRUCTURE (V)

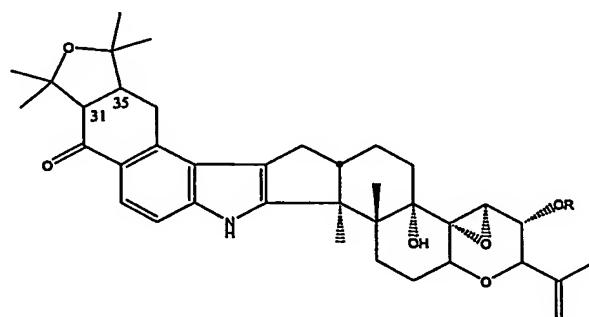
5 which includes compounds selected from the group consisting of: lolitriol; = $31\alpha, 35\beta$ stereochemistry where $R_1 = H$ or acetate and $R_2 = H$; lolitrem N = $31\alpha, 35\alpha$ stereochemistry where $R_1=H$ or acetate and $R_2=H$; Lolitrem J = $31\alpha, 35\beta$ stereochemistry where $R_1 = H$ or acetate and $R_2 = \text{acetate}$;



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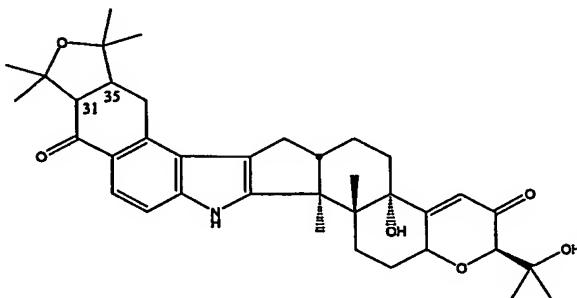
STRUCTURE (VI)

which includes lolitrem H = $31\alpha, 35\beta$ stereochemistry where $R = H$ or acetate;



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STRUCTURE (VII)

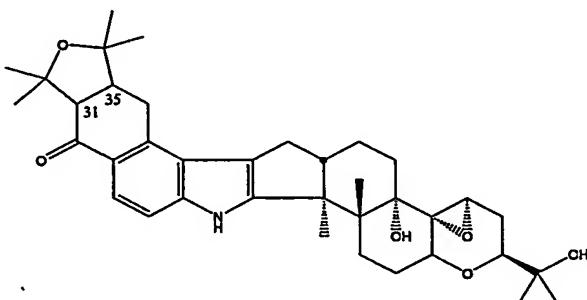
which includes lolitrem K = 31 α , 35 β stereochemistry, where R = H or acetate;



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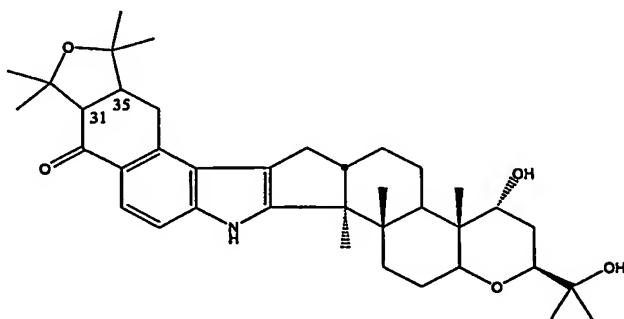
STRUCTURE (VIII)

which includes lolilline = 31 α , 35 β stereochemistry;



STRUCTURE (IX)

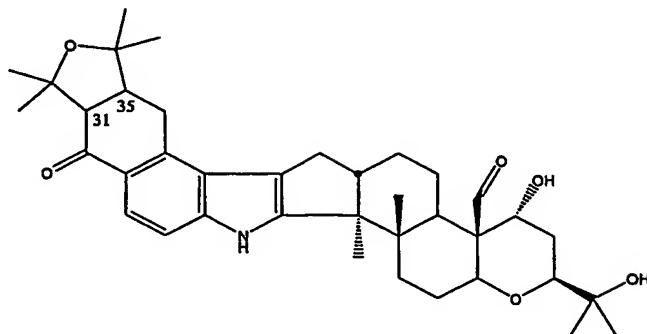
which includes lolitrem M = 31 α , 35 β stereochemistry;



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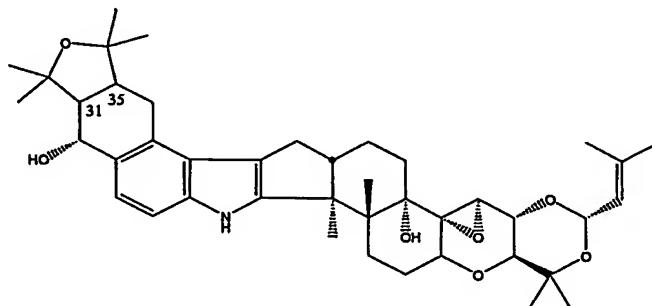
STRUCTURE (X)

which includes lolicine A = $31\alpha, 35\beta$ stereochemistry;



STRUCTURE (XI)

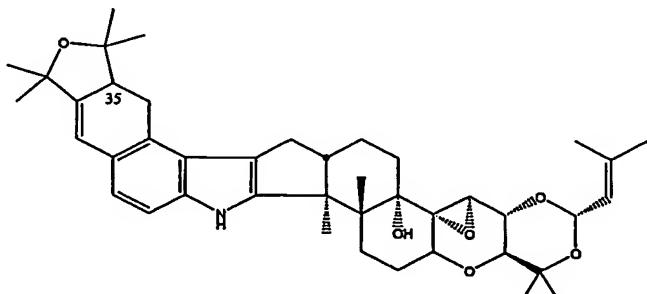
which includes lolicine B = $31\alpha, 35\beta$ stereochemistry;



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STRUCTURE (XII)

which includes compounds selected from the group consisting of: 30-desoxylolitrem B- 30α -ol = $31\alpha, 35\beta$ stereochemistry; 30-desoxy-31-*epi*lolitrem B- 30α -ol = $31\beta, 35\beta$ stereochemistry;



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STRUCTURE (XIII)

which includes 30-desoxyololitrem B-30-ene = 35β stereochemistry;

and combinations of the above compounds.

4. The method as claimed in any of the above claims wherein the composition further includes pharmaceutically and physiologically acceptable carriers.
5. The method as claimed in claim 4 wherein the pharmaceutically and physiologically acceptable carriers include components selected from the group including; fillers; excipients; modifiers; humectants; stabilisers; emulsifiers; diluents; and other formulation components such as a use of a lipid vehicle.
6. The method as claimed in any of the above claims wherein the composition is administered in a form selected from the group including: an injection; a tablet; a capsule; a suppository; an injection; a suspension; a drink or tonic; a syrup; a powder; an ingredient in solid or liquid foods; a nasal spray; a sublingual wafer; a transdermal patch; a transdermal injection; and combinations thereof.
7. The method as claimed in any of the above claims wherein the BK channel antagonist compound or compounds are extracted from endophyte-infected plants and seeds.
8. The method as claimed in any of claims 1 to 6 wherein the BK channel antagonist compound or compounds are extracted from fungal cultures.
9. The method as claimed in any of claims 1 to 6 wherein the BK channel antagonist compound or compounds are derived by chemical synthesis.
10. The method as claimed in any of claims 1 to 6 wherein the BK channel antagonist compound or compounds are extracted from heterologous expression systems including but not limited to bacteria, yeast, fungi, plants and animal cells.
11. The method as claimed in claim 7 wherein the perennial ryegrass seed is from

Lolium perenne.

12. The method as claimed in any of the above claims wherein the BK channel antagonist compound or compounds has activity against both alpha (α) subunit and alpha plus beta (β) accessory subunit (β_1 to β_4) channels.
- 5 13. The method as claimed in any of claims 1 to 4 wherein, for lolitrem B, the degree of antagonist inhibition is approximately 97% for a composition containing approximately 20nM lolitrem B.
- 10 14. The method as claimed in any of claims 1 to 4 wherein, for lolitrem B, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 3.7 ± 0.4 nM of lolitrem B.
- 15 15. The method as claimed in any of claims 1 to 4 wherein, for lolitriol, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 1000 nM lolitriol.
16. The method as claimed in any of claims 1 to 4 wherein, for lolitriol, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 195 nM of lolitriol to inhibit α and β_1 BK channel activity
17. The method as claimed in any of claims 1 to 4 wherein, for lolitriol, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 536 ± 16 nM of lolitriol to inhibit α and β_4 activity.
- 20 18. The method as claimed in any of claims 1 to 4 wherein, for 31-epilolitrem B, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 200nM 31-epilolitrem B.
19. The method as claimed in any of claims 1 to 4 wherein, for 31-epilolitrem B, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition

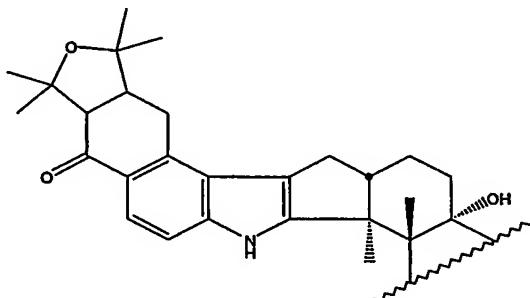
containing approximately 58 ±6 nM of 31-*epi*lolitrem B to inhibit α and β₁ activity.

20. The method as claimed in any of claims 1 to 4 wherein, for 31-*epi*lolitrem B, the half maximal degree of antagonist inhibition (IC₅₀) is found for a composition containing approximately 49 nM of 31-*epi*lolitrem B to inhibit α and β₄ activity.

5 21. The method as claimed in any of claims 1 to 4 wherein, for lolitrem E, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 100 nM lolitrem E.

22. The method as claimed in any of claims 1 to 4 wherein the antagonist effect of the composition is not able to be reversed by wash out for concentrations of 10 nM
10 or greater of lolitrem B compound.

23. Use of a composition for preventing repolarisation or hyperpolarisation of a cell that contains a BK channel wherein a pharmacologically effective amount of the composition is administered to the cell and wherein the composition contains at least one BK channel antagonist of the moiety shown in structure (I):



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STRUCTURE (I)

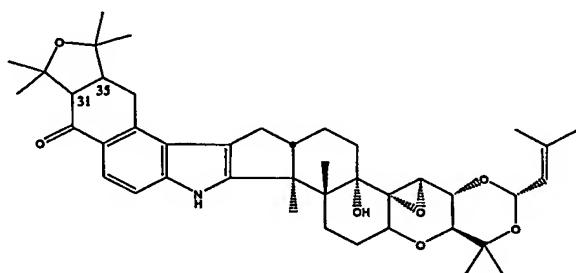
or derivatives thereof.

24. The use as claimed in claim 23 wherein the derivatives of structure (I) are selected from the group consisting of: salts, analogues, isomers, and combinations
20 thereof.

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25. The use as claimed in claim 23 or claim 24 wherein the antagonist compound is selected from the group consisting of: lolitrem B, lolitrem A, lolitrem F, 31-*epi*lolitrem F, 31-*epi*lolitrem B, lolitrem E, lolitrem E acetate, lolitrem L, lolitrem G, lolitrem C, lolitrem M, lolitriol, lolitriol acetate, lolitrem N, lolitrem J, lolitrem H,
5 lolitrem K, lolicine A and B, 30-desoxy lolitrem B-30 α -ol, 30-desoxy-31-*epi*lolitrem B-30 α -ol, 30-desoxylolitrem B-30-ene lolilline and combinations thereof.

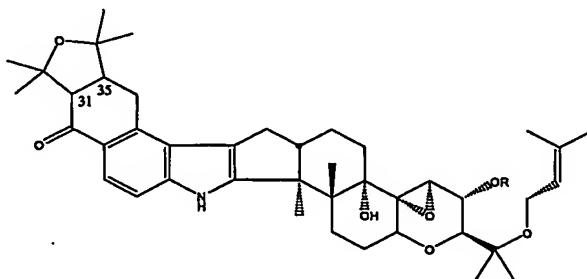
26. The use as claimed in claim 23 or claim 24 wherein the antagonist compound is selected from the group consisting of:



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STRUCTURE (II)

which includes compounds selected from the group consisting of: lolitrem B = 31 α , 35 β stereochemistry; 31-*epi*lolitrem B = 31 β , 35 β stereochemistry; lolitrem F = 31 α , 35 α ; 31-*epi*lolitrem F = 31 β , 35 α ;

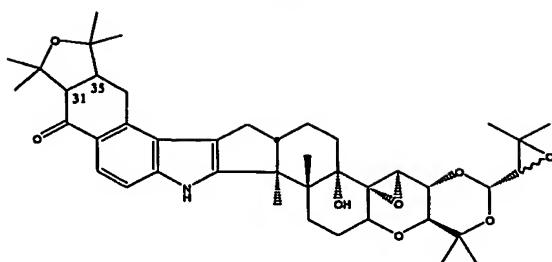


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STRUCTURE (III)

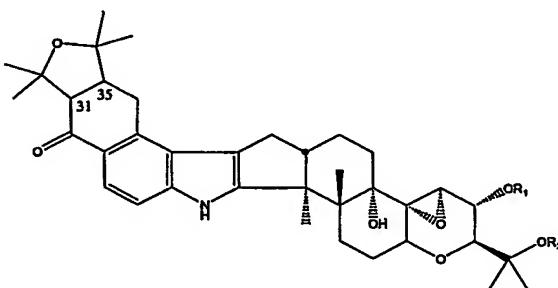
which includes compounds selected from the group consisting of: lolitrem E = 31 α , 35 β stereochemistry where R = H or acetate; lolitrem L = 31 α , 35 α stereochemistry where R = H or acetate;

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STRUCTURE (IV)

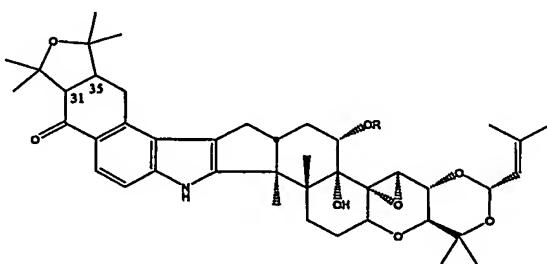
which includes compounds selected from the group consisting of: lolitrem A = $31\alpha, 35\beta$ stereochemistry; lolitrem G = $31\alpha, 35\alpha$ stereochemistry;



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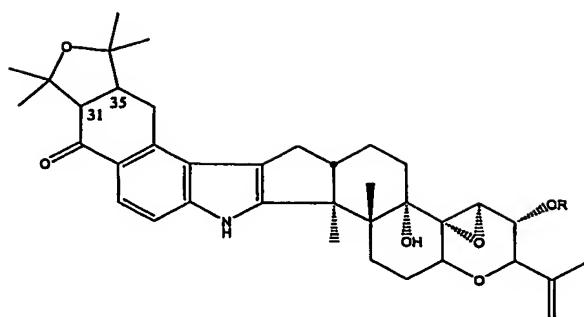
STRUCTURE (V)

which includes compounds selected from the group consisting of: lolitriol; = $31\alpha, 35\beta$ stereochemistry where $R_1 = H$ or acetate and $R_2 = H$; lolitrem N = $31\alpha, 35\alpha$ stereochemistry where $R_1=H$ or acetate and $R_2=H$; Lolitrem J = $31\alpha, 35\beta$ stereochemistry where $R_1 = H$ or acetate and $R_2 = acetate$;



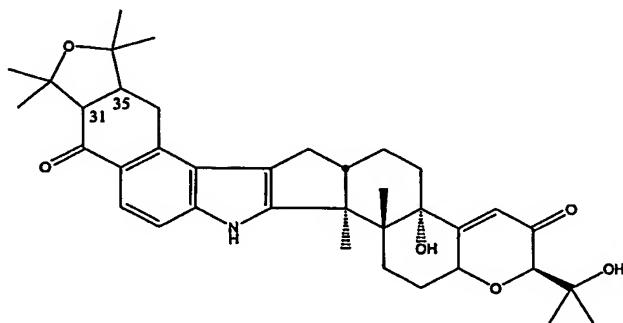
STRUCTURE (VI)

which includes lolitrem H = $31\alpha, 35\beta$ stereochemistry where $R = H$ or acetate;



STRUCTURE (VII)

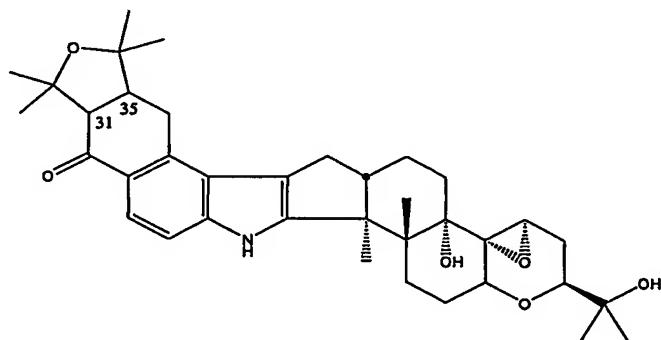
which includes lolitrem K = 31 α , 35 β stereochemistry, where R = H or acetate;



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STRUCTURE (VIII)

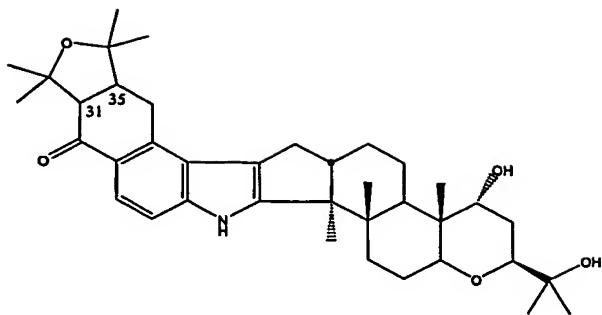
which includes lolilline = 31 α , 35 β stereochemistry;



STRUCTURE (IX)

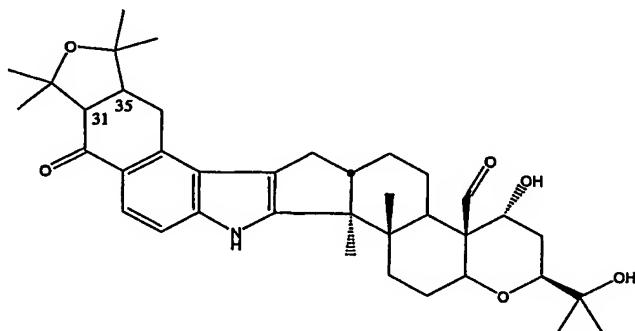
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which includes lolitrem M = 31 α , 35 β stereochemistry;



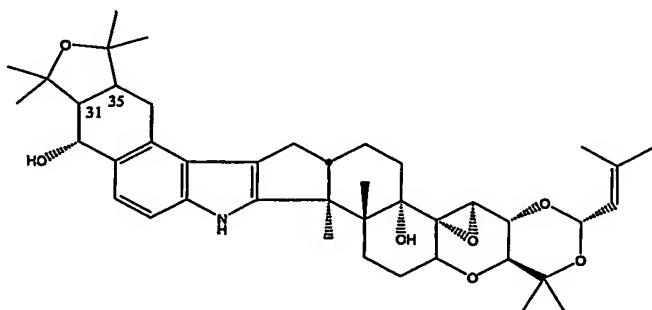
STRUCTURE (X)

which includes lolicine A = 31 α , 35 β stereochemistry;



STRUCTURE (XI)

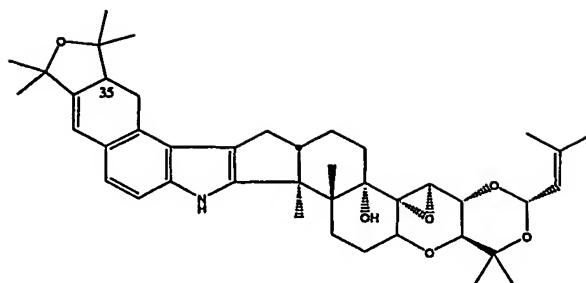
which includes lolicine B = 31 α , 35 β stereochemistry;



STRUCTURE (XII)

which includes compounds selected from the group consisting of:

- 10 desoxylolitrem B-30 α -ol = 31 α , 35 β stereochemistry; 30-desoxy-31-epiolitrem B-30 α -ol = 31 β , 35 β stereochemistry;



STRUCTURE (XIII)

which includes 30-desoxylolitrem B-30-ene = 35β stereochemistry;

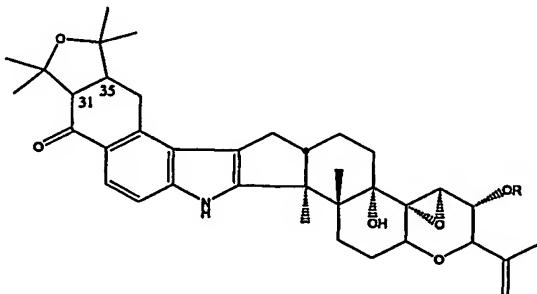
and combinations of the above compounds.

- 5 27. The use as claimed in any of the above claims wherein the composition further includes pharmaceutically and physiologically acceptable carriers.
28. The use as claimed in claim 27 wherein the pharmaceutically and physiologically acceptable carriers include components selected from the group including; fillers; excipients; modifiers; humectants; stabilisers; emulsifiers; diluents; 10 and other formulation components such as a use of a lipid vehicle.
29. The use as claimed in any of claims 23 to 28 wherein the composition is administered in a form selected from the group including: an injection; a tablet; a capsule; a suppository; an injection; a suspension; a drink or tonic; a syrup; a powder; an ingredient in solid or liquid foods; a nasal spray; a sublingual wafer; a 15 transdermal patch; a transdermal injection; and combinations thereof.
30. The use as claimed in any of claims 23 to 29 wherein the BK channel antagonist compound or compounds are extracted from endophyte-infected plants and seeds.
31. The use as claimed in any of claims 23 to 29 wherein the BK channel 20 antagonist compound or compounds are extracted from fungal cultures.

32. The use as claimed in any of claims 23 to 29 wherein the BK channel antagonist compound or compounds are derived by chemical synthesis.
33. The use as claimed in any of claims 23 to 29 wherein the BK channel antagonist compound or compounds are extracted from heterologous expression systems including but not limited to bacteria, yeast, fungi, plants and animal cells.
34. The use as claimed in claim 30 wherein the perennial ryegrass seed is from *Lolium perenne*.
35. The use as claimed in any of claims 23 to 34 wherein the BK channel antagonist compound or compounds has activity against both alpha (α) subunit and alpha plus beta (β) accessory subunit (β_1 to β_4) channels.
36. The use as claimed in any of claims 23 to 26 wherein, for lolitrem B, the degree of antagonist inhibition is approximately 97% for a composition containing approximately 20nM lolitrem B.
37. The use as claimed in any of claims 23 to 26 wherein, for lolitrem B, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 3.7 ± 0.4 nM of lolitrem B.
38. The use as claimed in any of claims 23 to 26 wherein, for lolitriol, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 1000 nM lolitriol.
39. The use as claimed in any of claims 23 to 26 wherein, for lolitriol, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 195 nM of lolitriol to inhibit α and β_1 BK channel activity
40. The use as claimed in any of claims 23 to 26 wherein, for lolitriol, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing

approximately 536 ± 16 nM of lolitrem B to inhibit α and β_4 activity.

41. The use as claimed in any of claims 23 to 26 wherein, for 31-*epi*lolitrem B, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 200nM 31-*epi*lolitrem B.
- 5 42. The use as claimed in any of claims 23 to 26 wherein, for 31-*epi*lolitrem B, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 58 ± 6 nM of 31-*epi*lolitrem B to inhibit α and β_1 activity.
43. The use as claimed in any of claims 23 to 26 wherein, for 31-*epi*lolitrem B, the half maximal degree of antagonist inhibition (IC_{50}) is found for a composition 10 containing approximately 49 nM of 31-*epi*lolitrem B to inhibit α and β_4 activity.
44. The use as claimed in any of claims 23 to 26 wherein, for lolitrem E, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 100 nM lolitrem E.
45. The use as claimed in any of claims 23 to 26 wherein the antagonist effect of 15 the composition is not able to be reversed by wash out for concentrations of 10 nM or greater of lolitrem B compound.
46. A composition that contains a pharmacologically effective amount of at least one BK channel antagonist compound containing the moiety shown in structure (VII):



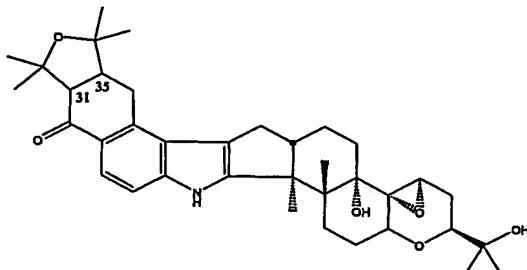
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STRUCTURE (VII)

which includes lolitrem K = $31\alpha, 35\beta$ stereochemistry, where R = H or acetate.

47. A composition that contains a pharmacologically effective amount of at least one BK channel antagonist compound containing the moiety shown in structure (IX):

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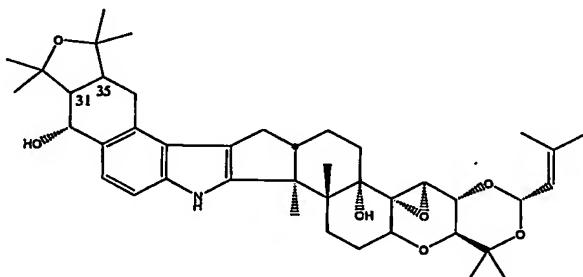


STRUCTURE (IX)

which includes lolitrem M = $31\alpha, 35\beta$ stereochemistry.

48. A composition that contains a pharmacologically effective amount of at least one BK channel antagonist compound containing the moiety shown in structure

10 (XII):

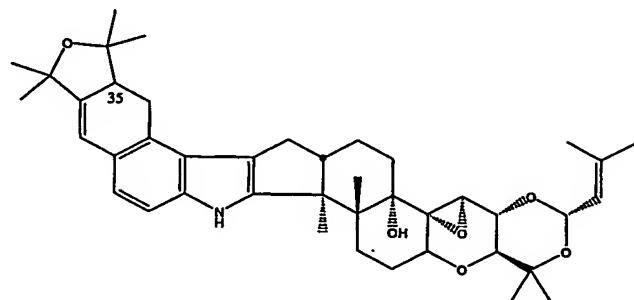


STRUCTURE (XII)

which includes compounds selected from the group consisting of: 30-desoxylolitrem B- 30α -ol = $31\alpha, 35\beta$ stereochemistry; 30-desoxy-31-*epi*lolitrem B- 30α -ol = $31\beta, 35\beta$ stereochemistry.

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49. A composition that contains a pharmacologically effective amount of at least one BK channel antagonist compound wherein the antagonist compound is structure (XIII):



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STRUCTURE (XIII)

which includes 30-desoxylolitrem B-30-ene = 35β stereochemistry.